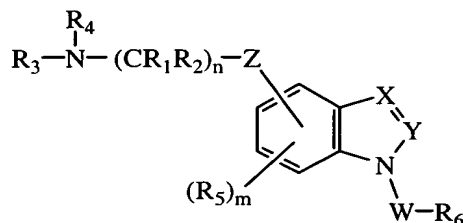


AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A compound of formula I



(I)

wherein

W is SO_2 , CO, CONH, CSNH or CH_2 ;

X is CR_7 or N ;

Y is CR_8 ;

Z is O, SO_p or NR_q ;

R_1 and R_2 are each independently H or C_1-C_6 alkyl;

n is an integer of 2, 3 or 4;

R_3 and R_4 are each independently H or a

cycloheteroalkyl ~~or heteroaryl~~ group each optionally substituted with the proviso that only one of R_3 or R_4 may be H, or R_3 and R_4 may be taken together with the atom to which they are attached to form an optionally substituted 3- to 6-membered ring containing an additional heteroatom selected from O, N or S;

R_5 is H, halogen, CN, OR_{13} , CO_2R_{14} , $CONR_{15}R_{16}$, $CNR_{17}NR_{18}R_{19}$,

$SO_2NR_{20}R_{21}$, SO_qR_{22} or a C_1-C_6 alkyl, C_2-C_6 alkenyl,

C_2-C_6 alkynyl, C_3-C_6 cycloalkyl, cycloheteroalkyl, or phenyl ~~or heteroaryl~~ group each optionally substituted;

m is an integer of 1, 2 or 3;

p and q are each independently 0 or an integer of 1 or 2;

R_6 is an optionally substituted C_1-C_6 alkyl, or aryl group;

R_7 and R_8 are each independently H, halogen or a C_1-C_6 alkyl, aryl, ~~heteroaryl~~ or C_1-C_6 alkoxy group each optionally substituted;

R_9 is H or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, cycloheteroalkyl, or aryl ~~or heteroaryl~~ group each optionally substituted;

R_{10} , R_{11} , R_{12} , R_{15} , R_{16} , R_{17} , R_{18} and R_{19} are each independently H or C_1 - C_4 alkyl;

R_{13} is H, COR_{23} , or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, or aryl ~~or heteroaryl~~ group each optionally substituted;

R_{14} is H or a C_1 - C_6 alkyl, or aryl ~~or heteroaryl~~ group each optionally substituted;

R_{20} and R_{21} are each independently H or a C_1 - C_6 alkyl, or aryl ~~or heteroaryl~~ group each optionally substituted; and

R_{22} and R_{23} are each independently an optionally substituted C_1 - C_6 alkyl, or aryl ~~or heteroaryl~~ group; or a pharmaceutically acceptable salt thereof.

2. (Original) The compound according to claim 1 wherein W is SO_2 .

3. (Original) The compound according to claim 1 wherein Z is O.

4. (Original) The compound according to claim 1 wherein n is 2.

5. (Previously Presented) The compound according to claim 1 wherein R_6 is an aryl group optionally substituted.

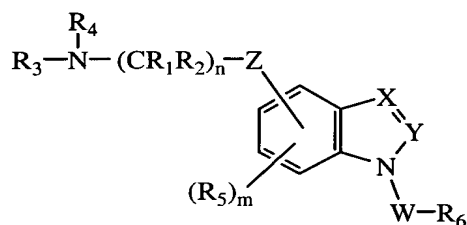
6. (Original) The compound according to claim 1 wherein X is CR_7 and R_5 and R_7 are H.

7. (Original) The compound according to claim 2 wherein R_1 and R_2 are H; Z is O; and n is 2.

8. (Currently Amended) The compound according to claim 6 wherein W is SO_2 ; Z is O; and R_3 and R_4 are taken together with the atom to which they are attached to form a 5- or 6-membered ring ~~optionally~~ containing one oxygen atom.

9. (Previously Presented) The compound according to claim 6 selected from the group consisting of:
4-(2-morpholin-4-ylethoxy)-1-(phenylsulfonyl)-1H-indole;
N-(2-{[1-(phenylsulfonyl)-1H-indol-4-yl]oxy}ethyl)tetrahydro-2H-pyran-4-amine; and
a pharmaceutically acceptable salt thereof.

10. (Withdrawn) A method for the treatment of a disorder of the central nervous system related to or affected by the 5-HT₆ receptor in a patient in need thereof which comprises providing to said patient a therapeutically effective amount of a compound of formula I.



(I)

wherein

W is SO₂, CO, CONH, CSNH or CH₂;

X is CR₇ or N;

Y is CR₈;

Z is O, SO_p or NR₉;

R₁ and R₂ are each independently H or C₁-C₆alkyl;

n is an integer of 2, 3 or 4;

R₃ and R₄ are each independently H or a cycloheteroalkyl, or heteroaryl group each optionally substituted with the proviso that only one of R₃ or R₄ may be H, or R₃ and R₄ may be taken together with the atom to which they are attached to form an optionally substituted 3- to 6-membered ring containing an additional heteroatom selected from O, N or S;

R₅ is H, halogen, CN, OR₁₃, CO₂R₁₄, CONR₁₅R₁₆, CNR₁₇NR₁₈R₁₉, SO₂NR₂₀R₂₁, SO_qR₂₂ or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, phenyl or heteroaryl group each optionally substituted;

m is an integer of 1, 2 or 3;

p and q are each independently 0 or an integer of 1 or 2;

R_6 is an optionally substituted C_1 - C_6 alkyl or aryl group;
 R_7 and R_8 are each independently H, halogen or a C_1 - C_6 alkyl, aryl, heteroaryl or C_1 - C_6 alkoxy group each optionally substituted;
 R_9 is H or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;
 R_{10} , R_{11} , R_{12} , R_{15} , R_{16} , R_{17} , R_{18} and R_{19} are each independently H or C_1 - C_4 alkyl;
 R_{13} is H, COR_{23} or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, aryl or heteroaryl group each optionally substituted;
 R_{14} is H or a C_1 - C_6 alkyl, aryl or heteroaryl group each optionally substituted;
 R_{20} and R_{21} are each independently H or a C_1 - C_6 alkyl, aryl or heteroaryl group each optionally substituted; and
 R_{22} and R_{23} are each independently an optionally substituted C_1 - C_6 alkyl, aryl or heteroaryl group; or
a pharmaceutically acceptable salt thereof.

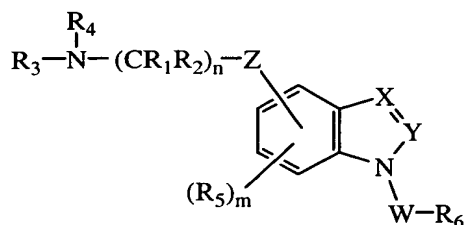
11. (Withdrawn) The method according to claim 10 wherein said disorder is a motor disorder, anxiety disorder or cognitive disorder.

12. (Withdrawn) The method according to claim 10 wherein said disorder is schizophrenia or depression.

13. (Withdrawn) The method according to claim 11 wherein said cognitive disorder is attention deficit disorder.

14. (Withdrawn) The method according to claim 11 wherein said cognitive disorder is Alzheimer's disease or Parkinson's disease.

15. (Currently Amended) A pharmaceutical composition which comprises a pharmaceutically acceptable carrier and an effective amount of a compound of formula I.



(I)

wherein

W is SO₂, CO, CONH, CSNH or CH₂;

X is CR₇ or N;

Y is CR₈;

Z is O, SO_p or NR₉;

R₁ and R₂ are each independently H or C₁-C₆alkyl;

n is an integer of 2, 3 or 4;

R₃ and R₄ are each independently H or a cycloheteroalkyl, ~~or heteroaryl~~ group each optionally substituted with the proviso that only one of R₃ or R₄ may be H, or R₃ and R₄ may be taken together with the atom to which they are attached to form an optionally substituted 3- to 6-membered ring containing an additional heteroatom selected from O, N or S;

R₅ is H, halogen, CN, OR₁₃, CO₂R₁₄, CONR₁₅R₁₆, CNR₁₇NR₁₈R₁₉, SO₂NR₂₀R₂₁, SO_qR₂₂ or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, or phenyl ~~or heteroaryl~~ group each optionally substituted;

m is an integer of 1, 2 or 3;

p and q are each independently 0 or an integer of 1 or 2;

R₆ is an optionally substituted C₁-C₆alkyl or aryl group;

R₇ and R₈ are each independently H, halogen or a C₁-C₆alkyl, aryl, ~~heteroaryl~~ or C₁-C₆alkoxy group each optionally substituted;

R₉ is H or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, or aryl ~~or heteroaryl~~ group each optionally substituted;

R₁₀, R₁₁, R₁₂, R₁₅, R₁₆, R₁₇, R₁₈ and R₁₉ are each independently H or C₁-C₄alkyl;

R₁₃ is H, COR₂₃ or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, or aryl ~~or heteroaryl~~ group each optionally substituted;

R₁₄ is H or a C₁-C₆alkyl, or aryl ~~or heteroaryl~~ group each optionally substituted;

R_{20} and R_{21} are each independently H or a C_1 - C_6 alkyl, or aryl ~~or heteroaryl~~ group each optionally substituted; and R_{22} and R_{23} are each independently an optionally substituted C_1 - C_6 alkyl, or aryl ~~or heteroaryl~~ group; or a pharmaceutically acceptable salt thereof.

16. (Original) The composition according to claim 15 wherein W is SO_2 ; Z is O; and n is 2.

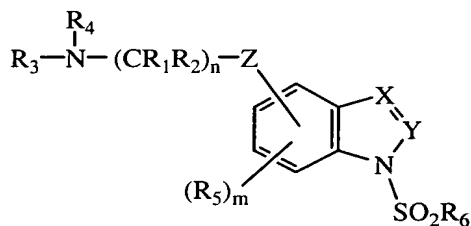
17. (Previously Presented) The composition according to claim 16 wherein R_6 is an aryl group optionally substituted.

18. (Original) The composition according to claim 17 wherein X is CR_7 , and R_1 , R_2 , R_5 , and R_7 are H.

19. (Previously Presented) The composition according to claim 18 having a formula I compound selected from the group consisting of:

4-(2-morpholin-4-ylethoxy)-1-(phenylsulfonyl)-1H-indole;
N-(2-{{1-(phenylsulfonyl)-1H-indol-4-yl}oxy}ethyl)tetrahydro-2H-pyran-4-amine; and
a pharmaceutically acceptable salt thereof.

20. (Withdrawn) A method for the preparation of a compound of formula Ia



(Ia)

wherein

X is CR_7 or N;

Y is CR_8 ;

Z is O, SO_p or NR_9 ;

R_1 and R_2 are each independently H or C_1 - C_6 alkyl;

n is an integer of 2, 3 or 4;

R₃ and R₄ are each independently H or a cycloheteroalkyl, or heteroaryl group each optionally substituted with the proviso that only one of R₃ or R₄ may be H, or R₃ and R₄ may be taken together with the atom to which they are attached to form an optionally substituted 3- to 6-membered ring containing an additional heteroatom selected from O, N or S;

R₅ is H, halogen, CN, OR₁₃, CO₂R₁₄, CONR₁₅R₁₆, CNR₁₇NR₁₈R₁₉, SO₂NR₂₀R₂₁, SO₂R₂₂ or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, phenyl or heteroaryl group each optionally substituted;

m is an integer of 1, 2 or 3;

p and q are each independently 0 or an integer of 1 or 2;

R₆ is an optionally substituted C₁-C₆alkyl or aryl group;

R₇ and R₈ are each independently H, halogen or a C₁-C₆alkyl, aryl, heteroaryl or C₁-C₆alkoxy group each optionally substituted;

R₉ is H or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

R₁₀, R₁₁, R₁₂, R₁₅, R₁₆, R₁₇, R₁₈ and R₁₉ are each independently H or C₁-C₄alkyl;

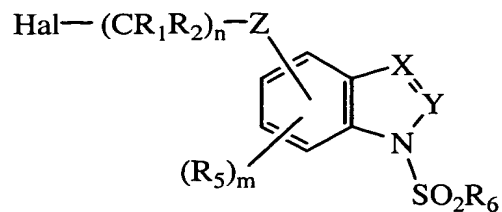
R₁₃ is H, COR₂₃ or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, aryl or heteroaryl group each optionally substituted;

R₁₄ is H or a C₁-C₆alkyl, aryl or heteroaryl group each optionally substituted;

R₂₀ and R₂₁ are each independently H or a C₁-C₆alkyl, aryl or heteroaryl group each optionally substituted; and

R₂₂ and R₂₃ are each independently an optionally substituted C₁-C₆alkyl, aryl or heteroaryl group

which method comprises reacting a compound of formula V'



(V')

wherein Hal is Cl, Br or I and X, Y, Z, n, m, R_1 , R_2 , R_5 and R_6 are as defined hereinabove with an amine, HNR_3R_4 , wherein R_3 and R_4 are defined hereinabove optionally in the presence of a solvent to give the desired compound of formula Ia.